AMENDMENTS TO THE CLAIMS

This listing of claims replaces all prior versions, and listings, of claims in the application.

1.-10. (Canceled)

11. (Currently Amended) A mutant ras peptide consisting of:

Xaa₁ Leu Xaa₂ Val Val Gly Ala Xaa₃ Gly Val Gly Lys Ser (SEQ ID NO:14);

wherein Xaa1 is the amino acid lysine or tyrosine;

wherein Xaa2 is an amino acid;

wherein Xaa₃ is selected from the group consisting of aspartic acid, valine, cysteine, alanine, arginine, and serine;

wherein when Xaa2 is valine, Xaa1 is tyrosine

and $\frac{1}{1}$ peptide elicits a peptide-specific human $CD8^+$ cytotoxic T lymphocyte immune response.

12. (Currently Amended) A mutant *ras* peptide <u>consisting of which is a fragment</u> of:

Xaa₁ Leu Xaa₂ Val Val Gly Ala Xaa₃ Gly Val Gly Lys Ser (SEQ ID NO:14) (SEQ ID NO: 15) or a fragment thereof,

wherein the peptide consists of 8 to 13 amino acids;

wherein the peptide includes Xaa₁;

wherein Xaa₁ is the amino acid lysine or tyrosine;

wherein Xaa2 is an amino acid;

wherein Xaa₃ is selected from the group consisting of aspartic acid, valine, cysteine, alanine, arginine, and serine;

wherein when Xaa2 is valine, Xaa1 is tyrosine;

wherein said peptide includes Xaa₁ and said the peptide elicits a peptide-specific human CD8⁺ cytotoxic T lymphocyte immune response.

13. (Currently Amended) A mutant *ras* peptide consisting of between 13 and 8 amino acids, wherein said peptide is the following peptide or a fragment thereof:

Xaa₁ Leu Xaa₂ Val Val Gly Ala Xaa₃ Gly Val Gly Lys Ser (SEQ ID NO:14) (SEQ ID

NO: 15) or a fragment thereof;

wherein the peptide consists of 8 to 13 amino acids;

wherein the peptide includes Xaa1;

wherein Xaa1 is the amino acid tyrosine;

wherein Xaa2 is valine;

wherein Xaa₃ is selected from the group consisting of aspartic acid, valine, cysteine, alanine, arginine, and serine;

wherein said peptide includes Xaa₁ and said the peptide elicits a peptide-specific human CD8⁺ cytotoxic T lymphocyte immune response.

14. (Currently Amended) A mutant *ras* peptide consisting of between 13 and 8 amino acids, wherein said peptide is the following peptide or a fragment thereof:

Xaa₁ Leu Xaa₂ Val Val Gly Ala Xaa₃ Gly Val Gly Lys Ser (SEQ ID NO:14) (SEQ ID NO: 15) or a fragment thereof;

wherein the peptide consists of 8 to 13 amino acids;

wherein the peptide includes Xaa1;

wherein Xaa1 is the amino acid lysine or tyrosine;

wherein Xaa₂ is selected from the group consisting of valine, tryptophan, leucine, tyrosine, and phenylalanine;

wherein Xaa₃ is selected from the group consisting of aspartic acid, valine, cysteine, alanine, arginine, and serine;

wherein when Xaa2 is valine, Xaa1 is tyrosine;

wherein said peptide includes Xaa₁ and said the peptide elicits a peptide-specific human CD8⁺ cytotoxic T lymphocyte immune response.

15. (Currently Amended) A mutant *ras* peptide consisting of between 13 and 8 amino acids, wherein said peptide is the following peptide or a fragment thereof:

Xaa₁ Leu Xaa₂ Val Val Gly Ala Xaa₃ Gly Val Gly Lys Ser (SEQ ID NO:14) (SEQ ID NO: 15) or a fragment thereof;

wherein the peptide consists of 8 to 13 amino acids;

wherein Xaa₁ is tyrosine;

wherein Xaa2 is an amino acid;

wherein Xaa3 aspartic acid;

and said the peptide elicits a peptide-specific human CD8⁺ cytotoxic T lymphocyte immune response.

16.-24. (Canceled)

25. (Previously Presented) A mutant *ras* peptide-carrier molecule conjugate comprising the mutant *ras* peptide consisting of Tyr Leu Val Val Val Gly Ala Asp Gly Val (SEQ ID NO:11) and a carrier molecule, wherein said carrier molecule enhances the immunogenicity of the peptide.

26. (Canceled)

27. (Previously Presented) An immunogen for eliciting a mutant *ras* peptide-specific human CD8⁺ cytotoxic T lymphocyte immune response comprising a mutant *ras* peptide of claim 72, wherein the immunogen elicits a mutant *ras* peptide-specific human CD8⁺ cytotoxic T lymphocyte immune response.

28.-31. (Canceled)

- 32. (Previously Presented) A pharmaceutical composition comprising the mutant *ras* peptide of claim 72 and a pharmaceutically acceptable carrier.
- 33. (Previously Presented) The pharmaceutical composition of claim 32, further comprising a biological response modifier.
- 34. (Previously Presented) The pharmaceutical composition of claim 32, further comprising a liposome formulation, an antigen presenting cell, or an adjuvant comprising mycobacterial cell wall skeleton and monophosphoryl lipid A.

35.-65. (Canceled)

66. (Previously Presented) A mutant *ras* peptide-carrier molecule conjugate comprising the mutant *ras* peptide consisting of Tyr Leu Val Val Val Gly Ala Asp Gly Val (SEQ ID NO:11) and a carrier molecule, wherein said carrier molecule enhances the immunogenicity of the peptide and wherein the carrier molecule is selected from the group

consisting of influenza peptide, tetanus toxoid-CD4 epitope, Pseudomonas exotoxin A, and poly-L-lysine.

- 67. (Previously Presented) A mutant *ras* peptide-carrier molecule conjugate comprising the mutant *ras* peptide consisting of Tyr Leu Val Val Val Gly Ala Asp Gly Val (SEQ ID NO:11) and a carrier molecule, wherein said carrier molecule enhances the immunogenicity of the peptide and wherein the carrier molecule is tetanus toxoid.
- 68. (Previously Presented) The pharmaceutical composition of claim 33, wherein the biological response modifier is interleukin 2.
 - 69. (Canceled)
- 70. (Previously Presented) The pharmaceutical composition of claim 32, further comprising interleukin 2, interleukin 6, interleukin 12, interferon, tumor necrosis factor, GM-CSF, β2-microglobulin, or combinations thereof.
- 71. (Previously Presented) The pharmaceutical composition of claim 33, further comprising a liposome formulation, an antigen presenting cell, or an adjuvant comprising mycobacterial cell wall skeleton and monophosphoryl lipid A.
- 72. (Previously Presented) A mutant *ras* peptide consisting of Tyr Leu Val Val Val Gly Ala Asp Gly Val (SEQ ID NO:11).